FORM PTO-1390 (REV. 11-2000) U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371

ATTORNEY 'S DOCKET NUMBER	U S. APPLICATION NO. (If known, see 37 CFR 1 5
UNVN.62457/05	09/856966

INTERNATIONAL APPLICATION NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED
PCT/US99/28038	24 November 1999	25 November 1999

TITLE OF INVENTION
BORON-CARBIDE SOLID STATE NEUTRON DETECTOR AND METHOD FOR USING THE SAME

APPLICANT(S) FOR DO/EO/US
DOWBEN, Peter A.; ADENWALLA, Shireen; ROBERTSON, Brian W.; BAI, Mengjun

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. [X] This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. [ ] This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. [ ] This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. [X] The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. [X] A copy of the International Application as filed (35 U.S.C. 371(c)(2))
   a. [ ] is attached hereto (required only if not communicated by the International Bureau).
   b. [ ] has been communicated by the International Bureau.
   c. [X] is not required, as the application was filed in the United States Receiving Office (RO/US).
6. [ ] An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
   a. [ ] is attached hereto.
   b. [ ] has been previously submitted under 35 U.S.C. 154(d)(4).
7. [X] Amendments to the claims of the International Aplication under PCT Article 19 (35 U.S.C. 371(c)(3))
   a. [ ] are attached hereto (required only if not communicated by the International Bureau).
   b. [ ] have been communicated by the International Bureau.
   c. [ ] have not been made; however, the time limit for making such amendments has NOT expired.
   d. [X] have not been made and will not be made.
8. [ ] An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).
9. [ ] An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. [ ] An English lanugage translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. [X] An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. [ ] An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. [X] A FIRST preliminary amendment.
14. [ ] A SECOND or SUBSEQUENT preliminary amendment.
15. [X] A substitute specification.
16. [ ] A change of power of attorney and/or address letter.
17. [ ] A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. [ ] A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. [ ] A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. [ ] Other items or information:

JC12 Rec'd PCT/PTO 25 MAY 2001

U.S. APPLICATION NO (if known, see 37 CFR 1.5) 09/856966	INTERNATIONAL APPLICATION NO PCT/US99/28038	ATTORNEY'S DOCKET NUMBER UNVN.62457/05

21. [X] The following fees are submitted:	CALCULATIONS PTO USE ONLY	
BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):		
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO **$1000.00**		
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO **$860.00**		
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO **$710.00**		
International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) **$690.00**		
International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) **$100.00**		
ENTER APPROPRIATE BASIC FEE AMOUNT =	$ 690	
Surcharge of **$130.00** for furnishing the oath or declaration later than [ ] 20 [X] 30 months from the earliest claimed priority date (37 CFR 1.492(e)).	$ 130	

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	13 - 20 =	0	x **$18.00**	$ 0	
Independent claims	3 - 3 =	0	x **$80.00**	$ 0	
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ **$270.00**	$ N/A	
TOTAL OF ABOVE CALCULATIONS =				$ 820	
[X] Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.			+	$ (410)	
SUBTOTAL =				$ 410	
Processing fee of **$130.00** for furnishing the English translation later than [ ] 20 [ ] 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				$ N/A	
TOTAL NATIONAL FEE =				$ 410	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). **$40.00** per property +				$ N/A	
TOTAL FEES ENCLOSED =				$ 410	
				Amount to be refunded:	$
				charged:	$

a. [X] A check in the amount of $ 410 to cover the above fees is enclosed.

b. [ ] Please charge my Deposit Account No. _____ in the amount of $ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.

c. [X] The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 19-2112. A duplicate copy of this sheet is enclosed.

d. [ ] Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. **Credit card information should not be included on this form.** Provide credit card information and authorization on PTO-2038.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

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Serial No.

International App. No. PCT/US99/28038

Filing Date:

BORON-CARBIDE SOLID STATE NUETRON DETECTOR AND METHOD FOR USING THE SAME

Attorney Docket No. UNVN. 62457

Examiner:

Art Unit:

CERTIFICATE OF MAILING
37 C.F.R. 1.8

I hereby certify that this correspondence is being deposited with the U.S. Postal Service as First Class Mail in an envelope addressed to: Assistant Commissioner for Patents, Washington, D.C. 20231, on:

5/25/2001
Date

Signature

PRELIMINARY AMENDMENT

Asst. Commissioner for Patents
Washington, DC 20231

Dear Sir:

The following preliminary amendment is being entered to ensure that the Patent Office is aware of the Article 34 amendments that were made in the applicants Response to Written Opinion filed on November 13, 2000 in the above-referenced application during Phase II of the Patent Cooperation Treaty. The Authorized Officer acknowledged the aforementioned amendments referring to them as Annexes in the Notification of Transmittal of International Preliminary Examination Report issued on December 22, 2000. A copy of the Amended Sheet and the original International Application are herein attached.

In the Specification

Page 1 has been amended by adding the number – 1 – to the top of the page.

Page 2, line 25 has been amended by deleting "Mev" and inserting "MeV".

Page 6, line 13 has been amended by deleting "(".

Page 6, line 26 has been amended by deleting "sec" and inserting – •s –.

Page 7, line 2 has been amended by deleting "mCurie" and inserting – mCi –.

Page 7, line 8 has been amended by deleting "microns" and inserting – nm –.

Page 7, line 11 has been amended by adding – nm – after "50".

Page 7, line 11 has been amended by deleting "microns" and inserting – nm –.

Page 7, line 12 has been amended by deleting "microns" and inserting – nanometers –.

Page 8, line 6 has been amended by deleting "micron" and inserting – nm –.

In the Claims

Claim 7, line 2, delete "microns" and insert – nm –.

Claim 11, line 2, delete "C" and insert – °C –.

Claim 12, line 3, delete "detector" and insert – neutron detecting device –.

Claim 13, line 3, "detector" and insert – neutron detecting device –.

In the Drawings

Figure 4, the vertical axis has been amended by deleting "per sec." and inserting – s^{-1} –.

Figure 5, the horizontal axis has been amended by deleting "NM" and inserting – nm –.

Figure 6, the horizontal axis has been amended by deleting "NM" and inserting – nm –.

Remarks

Examination of the application is requested.

Respectfully submitted,

Dennis B. Danella
Reg. No. 46,653

DBD

May 25, 2001

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THE COMMISSIONER IS HEREBY AUTHORIZED TO CHARGE ANY ADDITIONAL AMOUNT REQUIRED, OR CREDIT ANY OVERPAYMENT, TO ACCOUNT NO. 19-2112 A DUPLICATE COPY OF THIS SHEET IS ENCLOSED

BOORON-CARBIDE SOLID STATE NEUTRON DETECTOR AND METHOD OF USING SAME

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

The Board of Regents of the University of Nebraska acknowledges that some funding for the research leading to this application was provided by the United States Government.

CROSS REFERENCE TO RELATED APPLICATIONS

Not Applicable.

BACKGROUND OF THE INVENTION

The present invention relates to detection of neutrons. More specifically, the present invention relates to a method and device for the efficient detection of neutrons that employs a boron-rich semiconductor as an electrically active part of the detection device.

Neutron scattering is an important research method to determine the structure of solids and liquids. It is used to understand the forces that act between the atoms in these systems and to determine the magnetic behavior of materials as well. The research and practical applications cover a broad range of areas, from the basic properties of materials to studies of engineering and medical applications.

There are essentially only four elements suitable for forming solid state semiconductor neutron detectors – boron (B), cadmium (Cd), gadolinium (Gd) and lithium (Li). Lithium semiconductor materials exist ($LiInS_2$, $LiInSe_2$ and LiZnP) but are difficult to reliably fabricate into devices and are very difficult materials with which to work. Gadolinium conversion layer based silicon (Si) diodes have been fabricated and proposed for neutron detection, but are not particularly stable. Cadmium zinc telluride has been shown to yield thermal neutron detection and the cadmium neutron capture cross section is high, but the neutron capture produces such high energy gamma rays (over 0.5

MeV) that the detectors would have to be large in order to detect these gammas efficiently.

Use of boron with neutron detectors is known both in the scintillator, the gas and the conversion layer varieties. Boron phosphide (BP) heterojunction diodes with silicon were successfully tested as alpha radiation detectors, but failed to work as neutron detectors. Boron carbide (B_4C) was successfully used as a neutron detector based upon resistivity changes resulting from increased lithium doping, as were (111) BP wafers. The lithium production in the boron carbide was a result of the following nuclear reactions:

$$^{10}B + n \rightarrow {}^{7}Li\ (1.01MeV) + {}^{4}He\ (1.78MeV)$$

$$^{10}B + n \rightarrow {}^{7}Li\ (0.83\ MeV) + {}^{4}He\ (1.47\ MeV) + \gamma\ (0.48\ MeV)$$

Boron has also been considered as a coating to a silicon diode and a GaAs diode but the maximum efficiency is low (less than 5%).

Existing gas and liquid neutron detectors are much larger and less rugged than solid-state ones could be. However, existing solid state neutron detectors also suffer serious limitations. For example, known boron-doped semiconductors are only a few percent efficient because they contain relatively little boron. Gadolinium, lithium and hydrocarbon conversion layers are all adversely affected by corrosion and high temperatures.

Furthermore, known conversion layer devices have low efficiencies, unless multiply stacked, because the range of the reaction products in the material of the conversion layer is generally considerably less than the thickness required for stopping thermal neutrons. Gadolinium conversion layers are an exception - but the neutron - gadolinium reaction results in conversion electrons of relatively low energy (70 keV) compared with the reaction products in the case of neutron capture by boron 10. Cadmium zinc telluride has been shown to yield thermal neutron detection, but the neutron capture produces such high energy gamma rays (over 0.5 MeV) that the detectors must be large to detect these gammas efficiently. Scintillator combinations with photomultipliers or intensified cameras are bulky and heavy and, except for neutron-

detecting scintillating fibers coupled optically to a remote photomultiplier or camera, are intolerant of high temperatures.

Boron and boron compounds, including boron carbide, are also used in neutron absorbing shielding purposes in nuclear reactors and other types of neutron radiation environments. For example, boron carbide can be used with shielding, thermal electric power, or detection of neutrons (by means of the resistivity change not by detection of individual neutrons). However, use of boron carbide to detect neutrons where the boron carbide is an electrically active semiconductor is novel.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide an inexpensive solid state neutron detector that includes a robust, structurally forgiving boron rich semiconductor.

It is another object of the present invention to provide a boron carbide semiconductor that utilizes its electrical properties as a semiconductor rather than its electrical property of resistance as a means of detecting neutrons or its thermoelectric properties in detecting neutrons.

A still further object of the present invention is to provide a detection device that yields high gain.

A further object of the present invention is to provide a detection device that provides real time response.

A further object of the present invention is to provide a detection device that is capable of detecting single neutrons.

Yet another object of the present invention is to provide a detection device that has low sensitivity to gamma and other radiation.

Still another object of the present invention is to provide a method of detecting neutrons with a detector device having a boron carbide semiconductor.

According to the present invention, the foregoing and other objects are obtained by a detection device having a layer of boron carbide. In the device, the boron carbide layer is an electrically active part of the detection device. The sensing mechanism of the detection device is inherent in the electrically connected, semiconducting boron carbide layer, which provides neutron capture resulting in prompt,

innately highly amplified, electrical output signals following interception of neutron(s).

Additional objects, advantages, and novel features of the invention will be set forth in part in the description which follows, and in part will become apparent to those skilled in the practice of the invention. The objects and advantages of the invention may be realized and attained by means of the forms of instrument and the combinations particularly pointed out in the appended claims.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

In the accompanying drawings which form a part of the specification and which are to be read in conjunction therewith and in which like reference numerals are used to indicate like parts in the various views:

Fig. 1 is a schematic representation of a heterojunction diode embodying the present invention;

Fig. 2 is a schematic representation of the test device using the principles of the present invention.

Fig. 3 depicts voltage-current characteristics of heterojunction diodes of the present invention;

Fig. 4 depicts count rates of neutrons with insertion of heterojunction diodes of the present invention into a neutron reactor; and

Figs. 5 and 6 depict the relationship of ideally attainable neutron detection efficiency as a function of the thickness of the boron-carbide layer of heterojunction diodes of the present invention in the cases of natural 10 Boron abundance and 100% 10 Boron enrichment of the boron carbide layer.

DETAILED DESCRIPTION OF THE INVENTION

Referring initially to Fig. 1, a heterojunction diode 10 is shown. This invention also applies to homojunction diodes and other known semiconductor detection devices, examples of which are provided below. Diode 10 is shown as having a boron carbide boron-carbon alloy semiconductor 12 on a silicon substrate 14. Semiconductor 12 is grown by plasma-enhanced chemical vapor deposition (PECVD). The preferred deposition technique is disclosed in U.S. Patent Nos. 4,957,773 (Spenser, et al.); 5,468,978 (Dowben); 5,658,834 (Dowben), which patents are expressly incorporated by reference herein. A pair of sputter-deposited gold electrodes 16 communicate with

semiconductor 12 and substrate 14. Secured to each electrode 16 is a wire 18 that serves to connect electrodes 16 to a bias voltage source and an electrical detection device such as a charge pulse measurement circuit. The sensory/measurement devices as well as monitoring devices are known and will not be discussed further.

Essentially, the invention works by including a boron-rich carbon alloy as an electrically active semiconductor region of a detector and by placing the detector where it can receive neutrons. The preferred way to detect neutrons is with atoms which are the most likely to capture neutrons and in which each neutron capture leads to the creation of one or more energetic charged particles whose mass is large compared with that of an electron and whose energy is large and can efficiently be converted to a measurable electrical signal. Boron atoms are highly likely to capture neutrons and such neutron capture creates highly energetic ions.

The following two reactions between boron isotope 10 (^{10}B) and a thermal neutron form the basis for neutron detection as contemplated by the present invention:

$$^{10}B+n_{thermal} \rightarrow {}^{7}Li\ (0.84MeV) + {}^{4}He\ (1.47MeV) + \gamma\ (0.48MeV)$$

$$^{10}B + n_{thermal} \rightarrow {}^{7}Li\ (1.01MeV) + {}^{4}He\ (1.78MeV)$$

With a boron-rich semiconductor, the boron captures the neutron and promptly decays into high-energy ions. The energetic ions cause secondary ionization of orders of magnitude more atoms in the surrounding materials for each captured neutron, liberating a correspondingly large electrical charge. The diode nature of the device enables the electrical charge to be collected. Also, incorporating the boron-rich alloy as an electrically active semiconductor part of the detector allows for the overall thickness of the device to be reduced while retaining high efficiency of neutron detection.

The first device to use this concept was a boron-carbon alloy semiconductor (grown by plasma-enhanced chemical vapor deposition) on a silicon substrate with sputter-deposited gold electrodes, as shown in Fig. 2. As seen in Fig. 2, a boron carbide/silicon diode 20 is connected to a charge sensitive preamplifier 22.

Charge sensitive preamplifier 22, in turn, is connected to a bias voltage input 24 and a single channel analyzer / multichannel scaler 26 which is connected to a computer 28.

In this heterojunction diode, the above reactions lead to dense local ionization of atoms and hence production of electron-hole pairs (at least of order $5x10^5$ pairs per neutron reaction), many of which are collected due to the applied bias voltage and form a charge pulse which is registered and counted by external circuitry. Such a device was first tested successfully on July 24, 1998 at the nuclear reactor in the VA Hospital in Omaha, Nebraska. This device could be improved in several ways, including ^{10}B-enrichment (to nearly 100% ^{10}B from the naturally occurring approximately 19% ^{10}B found in unenriched boron), increasing the thickness and quality of the boron carbide layer, changes in the electrical configuration and electrical circuitry, and changes in the functional and geometrical configurations.

The deposition of films for the heterojunction diodes (boron-carbon alloy, B_5C, on (111) Si) performed in this test was undertaken in a custom designed parallel plate 13.56 (MHz radio-frequency PECVD reactor used in previous studies). The silicon substrates were doped to $7x10^{14}/cm^3$. The (111) Si substrates surfaces were prepared by Ar+ ion sputtering in the plasma reactor. The source molecule gas closo-1,2-dicarbadodecaborane (ortho-carborane, $C_2B_{10}H_{12}$) was used as the source compound for growing the boron carbon alloy.

Typical B_5C/n-type silicon heterojunctions have been routinely formed by this technique. An example of one such diode device is presented in Fig. 2 with the boron carbide alloy layer of about 1000 nm thick as used as a neutron detector. These devices typically have onsets of 1 eV with very little leakage current (less than 5 μA at 25°C) and the boron carbide layer has the p-type character of the undoped PECVD semiconducting boron carbide in this device topology.

The detector area of these heterojunction diodes was about 1 cm^2, and wired in a "mesa" geometry. The neutron source was a small TRIGA-type reactor (V.A. Medical Center, Omaha, NE) with a flux of $1.6x10^6$ $n/cm^2 \cdot s$ based on calculations for the fission chamber. A heterojunction diode, reversed biased to about 3 V, was wired for pulse counting as shown in Figure 2 and inserted into the reactor. The resulting count

rates with insertion are plotted in Figure 3. Background and noise counts are in the range of 250 to 300 Hz, and within the reactor, the count rate rises to $2x10^5$ Hz.

To assure that very little of this count rate is attributable to gamma radiation, the diode was tested against a 100 mCi ^{137}Cs source for gamma radiation at a distance of 10 cm. The 661 keV gamma rays provided no detectable increase in count rate above background in spite of an expected 10^6 gamma rays incident on the diode per second. This is consistent with the expected extremely low gamma-ray sensitivity of such a solid state boron-carbon/silicon semiconductor alloy device, since boron and carbon have low atomic numbers and the boron-rich detectors were made very thin (1000 nm), and the electrically active silicon layer was under 600 nm thick.

Given that almost all counts are attributable to neutrons and that the boron carbide film is about 1000 nm thick, the detection efficiency is thus about 1%. Given that devices can be made with boron carbide of 50 nm to 100 nm in thickness and with depletion layers extending several nanometers, the single (thermal) neutron detection efficiencies are, conservatively, expected to reach 80% and higher in devices which simultaneously have exceedingly low γ-ray sensitivity (< 1% detection efficiency for all energies greater than 100 keV and < 0.01% for all energies above 0.5 MeV, assured by the use of boron as the dominant atomic species). Since the neutron - ^{10}B interaction results almost exclusively in the yield of highly ionizing lithium ions and alpha particles of total kinetic energy about 1.5 MeV and the boron atoms form the major species in the active semiconducting regions of the devices, the boron-carbon alloy layer of the detector yields an enormous internal gain (considerably greater than 10^5) which is essentially noise-free and comparable with the gain of the intensifiers and photomultipliers commonly used in scintillation-based detectors and imagers. By using exclusively ^{10}B enriched boranes in the PECVD fabrication process, detection efficiency with thinner films can be considerably improved compared with devices whose ^{10}B content reflects the natural isotopic abundance, about 19% ^{10}B.

As seen in Figure 2, the electronics demands are minimal compared with those for gadolinium neutron conversion layer-based detectors (which rely on the much smaller 70 keV energetically available for signal generation by the conversion electrons from gadolinium), while ensuring considerably greater efficiency and stability.

Additionally, the boron-carbon devices can be thinner than 100 μm thick and still achieve nearly 100% thermal neutron detective efficiency. Stacking diodes, interleaved with neutron energy absorbers, to form efficient neutron "calorimeters" or spectrometers is also possible. In combination with boron carbide based high temperature electronics, the boron-carbon based neutron detection systems are expected to be particularly applicable in harsh environments because of the refractory and mechanical performance of boron carbide. The boron-carbon devices may even be fabricated on metal substrates as well as fabricated with spatial resolution that could be on scales smaller than 0.5 nm. There is the possibility of fabricating spatial array detectors, including position sensors for scattering experiments, as well.

High efficiency is achieved because there is a proportionally large amount of boron present in the semiconductor layer. The boron carbide semiconductor has boron of whatever isotope one therefore chooses present in atomic fractions in the order of 80%. This is exceedingly rich in boron compared with any other suitable semiconductor. Because the density of boron atoms in the material is so high, the boron-rich layer can be quite thin and still contain enough boron atoms per unit area to be able to detect the neutrons very efficiently. In naturally occurring boron there is close to 20% of the boron atoms which are ^{10}B atoms which are the isotopes which interact strongly with neutrons to give the reactions provided above. It is certainly possible to increase the fraction of boron that is ^{10}B from natural abundance to about 95% or higher. This enrichment would result in ^{10}B atoms accounting for a fraction, about 80% or higher, of all atoms in the semiconductor boron carbide layer. Hence, if material enriched in ^{10}B is used rather than just the naturally occurring isotope ratio of ^{10}B, the efficiency increases even further.

Another important issue for efficiency is not just the reaction of the neutron with boron, but the ability to detect the reaction. By incorporating the boron atoms in an electrically active semiconductor where the lithium ion and the alpha particle can cause dense ionization of other atoms, many electron-hole pairs can be created by ionization of the atoms, and the electric fields that can be applied across the boron carbide layer can sweep out a large fraction of the electron-hole pairs. Thus, there are three aspects to efficiency. The first is ^{10}B being present in large number density. The second

being that the reaction of ^{10}B with neutrons results in ions which very efficiently ionize atoms in the surrounding in an electrically active semiconductor where the charge can be swept out efficiently. The third aspect of efficiency is that ^{10}B results in ions which have such a large energy that they can produce very large numbers of detectable electron hole pairs. The reactions which occur between neutrons and the other elements which give probable neutron interactions don't result in reaction products which are as readily detectable or detectable to give such large signals. Boron is unique.

Another point concerns detection devices having conversion layers containing boron. Neutron capture by boron generates the alpha particle and the lithium ion which can only travel a very limited distance. If conversion layer contained enough boron atoms to cause capture of a sufficient fraction of neutrons, then the layer will be so thick that the lithium and the alpha particles in some cases will not get out of the boron layer and, therefore, will not generate signals that are readily detectable. This is a severe defect compared with the boron carbide semiconductor devices of the present invention.

This invention can be used in various forms of solid-state neutron detectors presenting entrance detecting areas of order μm^2 to m^2. These detectors are capable of being implemented with very thin detecting and electrically active regions (≤ 1 μm minimum effective electrical thickness), with very low mass per unit detecting area, with efficiencies ranging up to nearly 100% even for single neutrons, with real-time response, with high spatial resolution (≤ 1 μm minimum), and with high temporal resolution. Of course, implementation may not always need to, or be able to, employ each of these attributes. Voltage and power needs are slight, as are charge pulse processing requirements.

Although the invention is described above as relating to heterojunction diodes, it is to be understood that the invention can be implemented in a large number of other ways, including homojunction diodes; p-i-n diodes; metal-semiconductor-metal, Schottky and other diodes; transistors; diode and transistor arrays; charge-induced devices (CID) and CID arrays; charge-coupled devices (CCD) and CCD arrays; solid-state neutron-detecting analogs of "photomultipliers"; neutron semiconductor avalanche devices; position-sensitive detectors, including those relying on charge subdivision or sensing and on current subdivision and those having capacitive or resistive means of

doing so; semiconductor drift detectors or semiconductor drift chambers; stacked series of one or more of the above detector types which are configured to serve as neutron energy spectrometers; individual or stacked series of one or more of the above detector types which also, or alternatively, serve as dosimeters. The dosimeters can be capable of yielding both real-time and cumulative dosimetry information once or many times, completely nondestructively of the dosimetry information contained in the detectors.

The range of applicability of the present invention includes: medical radiation dosimetry; detecting nuclear material; anti-terrorism and anti-smuggling devices; monitoring of nuclear reactors, of nuclear storage units and facilities, and of nuclear weapons, weapons storage and weapons shipment; life science materials and physical sciences scattering experiments; monitoring of neutron sources; calibration of neutron flux; personnel and environmental radiation protection; radiation protection at high energy radiation facilities, including medical x-ray facilities (high energy ones); neutron cancer therapy; profiling of medical, therapeutic, research and other neutron beams; comet, planetary and other space exploration.

From the foregoing, it will be seen that this invention is one well adapted to attain all the ends and objects herein above set forth together with other advantages which are obvious and which are inherent to the structure. It will be understood that certain features and subcombinations are of utility and may be employed without reference to other features and subcombinations. This is contemplated by and is within the scope of the claims.

Since many possible embodiments may be made of the invention without departing from the scope thereof, it is to be understood that all matter herein set forth or shown in the accompanying drawings is to be interpreted as illustrative and not in a limiting sense.

Having thus described the invention, what is claimed is:

1. A neutron detection device, said device comprising: a sensing mechanism, said sensing mechanism having a layer of boron carbide semiconductor wherein the boron carbide layer is an electrically active part of said detection device; and a monitoring device, wherein said monitoring device records said changes in said boron carbide layer detected by said sensing mechanism.

2. The device of claim 1, wherein said sensing mechanism is inherent in said boron carbide semiconductor layer and results in a prompt, innately highly amplified, electrical output following capture of a single neutron.

3. The device of claim 2, wherein said device is a homojunction diode.

4. The device of claim 1, further comprising a layer of silicon communicating with said layer of boron carbide.

5. The device of claim 4, wherein said device is a heterojunction diode.

6. The device of claim 1, wherein the thickness of said boron carbide layer is about 1000 nm.

7. The device of claim 5, wherein the thickness of said silicon layer is less than 600 nm.

8. The device of claim 1, further comprising at least two diodes interleaved with a neutron energy absorber.

9. The device of claim 1, wherein said boron carbide layer is fabricated on a metal substrate.

10. The device of claim 1, wherein said boron carbide layer contains at least 80% ^{10}B.

11. The device of claim 1, wherein said device is capable of operating at 500 °C.

12. A method of detecting neutrons, said method comprising: positioning a neutron detecting device in a location to allow said device to intercept a stream of neutrons, said detector comprising a layer of boron carbide wherein said boron carbide layer is an electrically active part of said device, and a sensing mechanism coupled to said boron carbide layer; introducing at least one neutron traveling in a direction to be intercepted by the boron carbide layer; and monitoring the interaction of the neutron with

the boron carbide semiconductor; wherein said sensing mechanism detects changes in said boron carbide layer caused by the interception of neutrons.

13. A method of detecting neutrons, said method comprising: positioning a neutron detecting device in a location to allow said device to intercept a stream of neutrons, said detector comprising a layer of boron carbide wherein said boron carbide layer is an electrically active part of said device, and a sensing mechanism inherent to said boron carbide layer; introducing at least one neutron traveling in a direction to be intercepted by the boron carbide layer; and monitoring the interaction of the neutron with the boron carbide semiconductor; wherein said sensing mechanism detects changes in said boron carbide layer caused by the interception of neutrons.

ABSTRACT OF THE DISCLOSURE

A boron carbide solid state neutron detector and method of using the detector is disclosed, wherein the detector includes a layer of boron carbide wherein the boron carbide layer is an electrically active part of the detection device, a sensing mechanism inherent to said boron carbide layer, wherein the sensing mechanism detects changes in the boron carbide layer caused by the interception of neutrons and a monitoring device coupled to the sensing mechanism.

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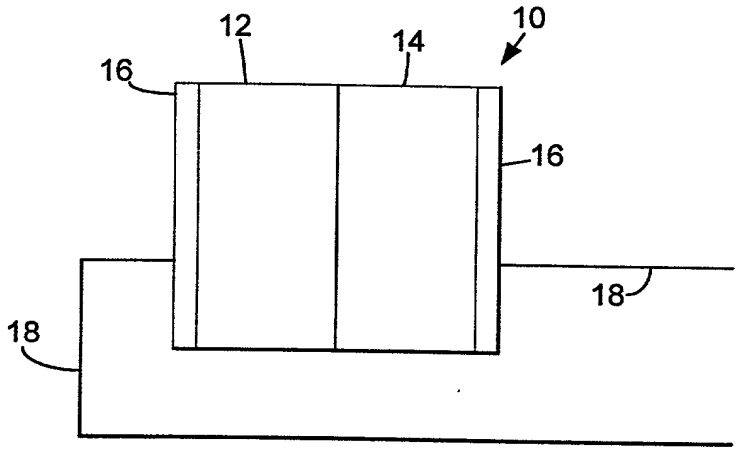


FIG.1.

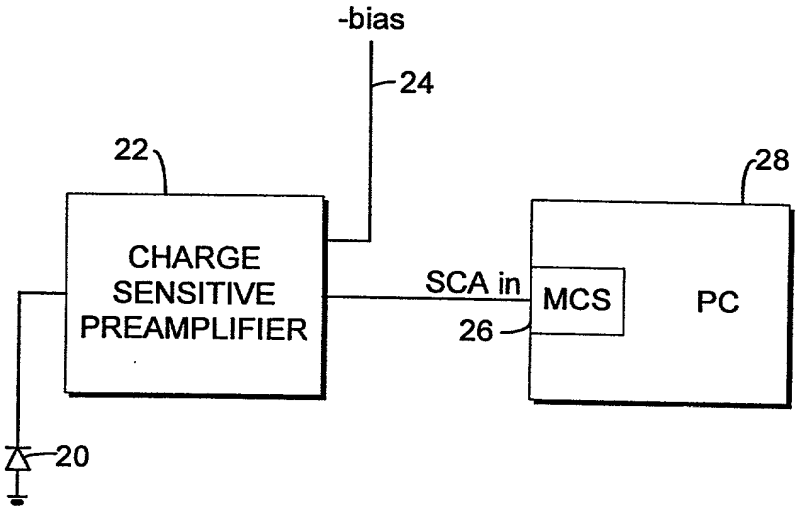
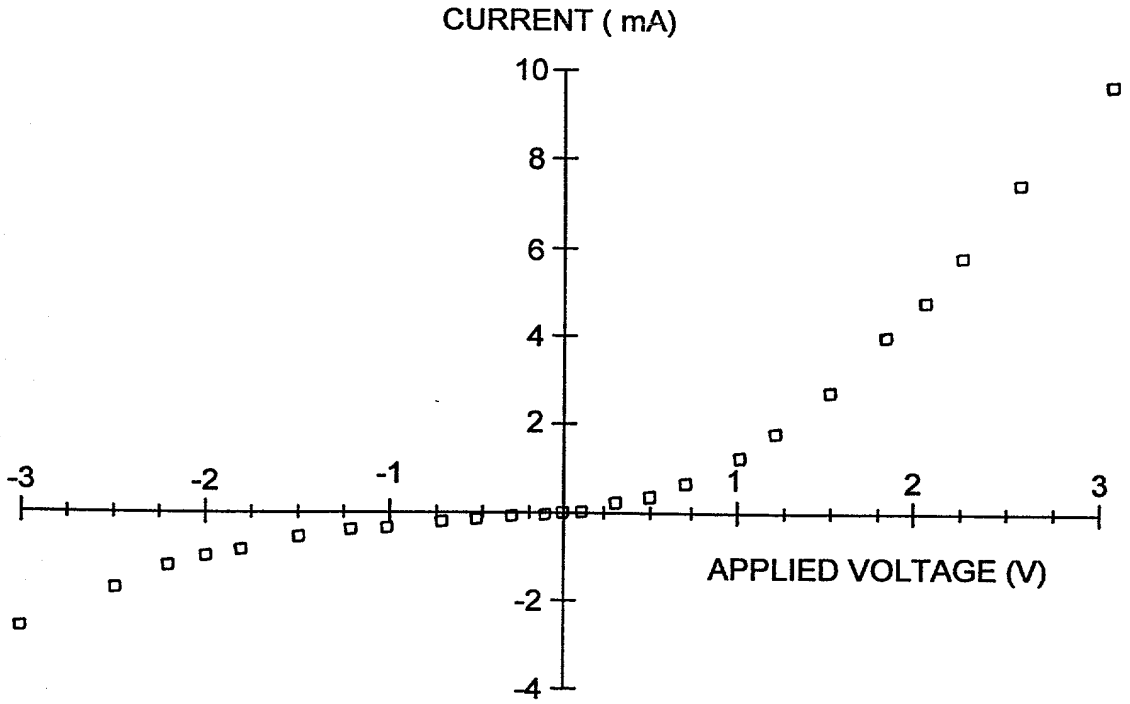

-bias
24
22
28
CHARGE
SENSITIVE
PREAMPLIFIER
SCA in
MCS
PC
26
20

FIG.2.

FIG.3.



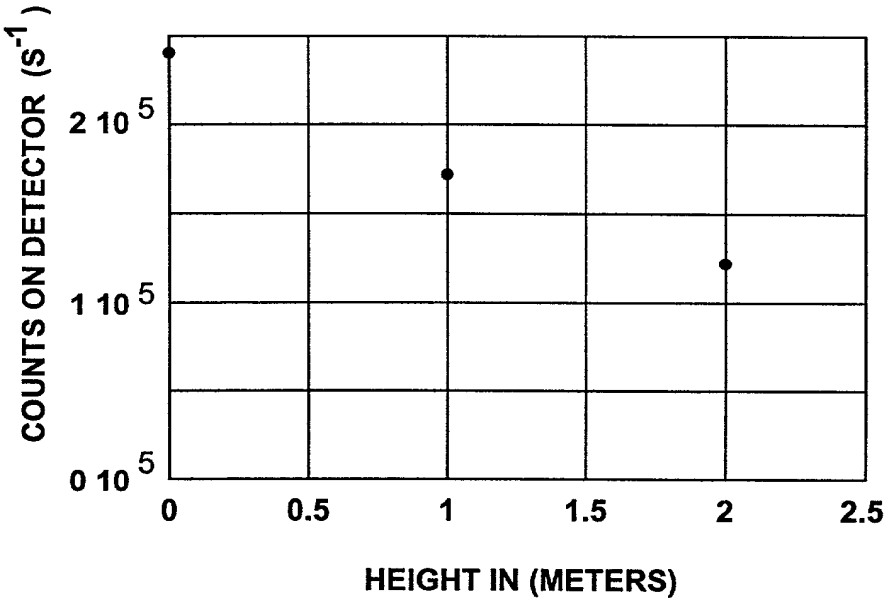
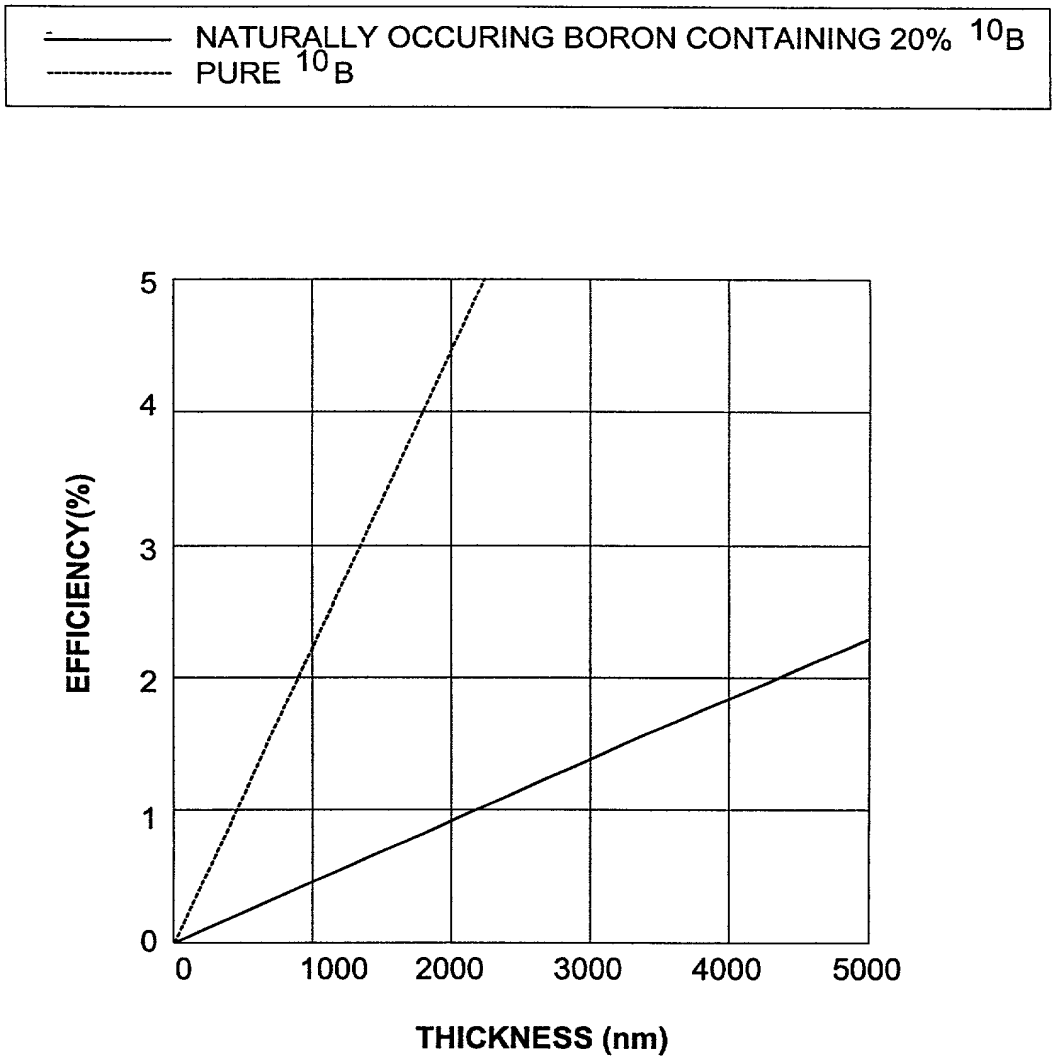


FIG. 4.

FIG. 5.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PURE 10B
EFFICIENCY(%)
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2
3
4
5
0
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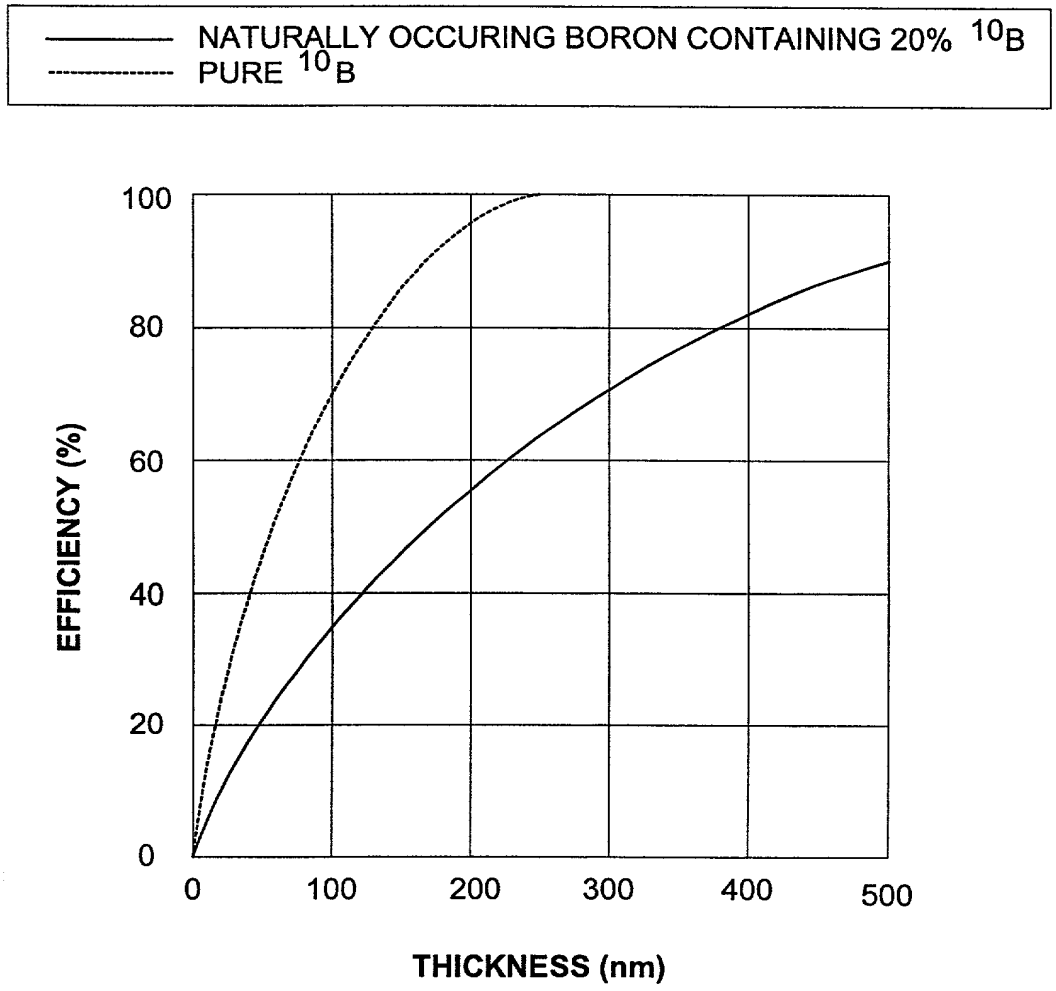
2000
3000
4000
5000
THICKNESS (nm)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PURE 10B
EFFICIENCY (%)
0
20
40
60
80
100
0
100
200
300
400
500
THICKNESS (nm)

FIG. 6.

JOINT DECLARATION FOR PATENT APPLICATION

As below-named inventors, we hereby declare that:

Our residence, post office address and citizenship are as stated below next to our respective names.

We believe we are the original, first and joint inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled BORON-CARBIDE SOLID STATE NEUTRON DETECTOR AND METHOD FOR USING THE SAME, the specification of which was filed on May 25, 2001, as Application Serial No. 09/856,966.

We hereby state that we have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, 1.56(a).

We hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Claimed? Y/N	Certified Copy Attached? Y/N

We hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below:

Application Number(s)	Filing Date (MM/DD/YYYY)
60/109,898	25 Nov. 1998

We hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, we acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

Application Number(s)	Filing Date (MM/DD/YYYY)
PCT/US99/28038	24 Nov. 1999

We hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith and to file and prosecute any corresponding foreign applications, including any international applications under the Patent Cooperation Treaty or the European Patent Convention: William B. Kircher, Reg. No. 22,481; James H. Marsh, Jr., Reg. No. 24,533; J. David Wharton, Reg. No. 25,717; Joseph B. Bowman, Reg. No. 25,807; Peter W. Gowdey, Reg. No. 25,872; Richard R. Johnson, Reg. No. 27,452; Walter R. Brookhart, Reg. No. 29,518; James H. Riley, II, Reg. No. 31,131; Joan Optican Herman, Reg. No. 31,968; Michael B. Hurd, Reg. No. 32,241; Michael J. Gross, Reg. No. 35,528; William P. Jensen, Reg. No. 36,833; Chris Murphy, Reg. No. 39,786; Daniel W. Shinn, Reg. No. 40,810; B. Trent Webb, Reg. No. 40,865; Susan J. Wharton, Reg. No. 41,524; Scott B. Strohm, Reg. No. 42,172; Janine A. Carlan, Reg. No. 42,387; Clinton G. Newton, Reg. No. 42,930; Ladi Shogbamimu, Reg. No. 46,291; Dennis Danella, Reg. No. 46,653; Tawni L. Brown, Reg. No. 47,456; Daniel P. Devers, Reg. No. 47,523; and Greg S. Donahue, Reg. No. 47,531. Address all correspondence to: Michael J. Gross, SHOOK, HARDY & BACON L.L.P., One Kansas City Place, 1200 Main Street, Kansas City, Missouri 64105-2118, telephone number (816) 474-6550.

We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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